

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : D21H 17/67, 17/64		A1	(11) International Publication Number: WO 99/49133 (43) International Publication Date: 30 September 1999 (30.09.99)
<p>(21) International Application Number: PCT/CA99/00213 (22) International Filing Date: 17 March 1999 (17.03.99) (30) Priority Data: 60/079,097 23 March 1998 (23.03.98) US (71) Applicant: PULP AND PAPER RESEARCH INSTITUTE OF CANADA [CA/CA]; 570 Boulevard St-Jean, Pointe Claire, Quebec H9R 3J9 (CA). (72) Inventors: MIDDLETON, Steven, R.; Apartment 703, 217 Stillview Avenue, Pointe Claire, Quebec H9R 2Y5 (CA). DESMEULES, Josee; 5034 rue Adam, Montreal, Quebec H1V 1W6 (CA). SCALLAN, Anthony, M.; 162 Coolbreeze, Pointe Claire, Quebec H9R 3S7 (CA). (74) Agents: MURPHY, Kevin, P. et al.; Swabey Ogilvy Renault Suite 1600, 1981 McGill College Avenue, Montreal, Quebec H3A 2Y3 (CA).</p> <p>(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p>			
<p>Published <i>With international search report.</i></p>			

(54) Title: METHOD FOR PRODUCING PULP AND PAPER WITH CALCIUM CARBONATE FILLER

(57) Abstract

A method is described for attaining high levels of loading of calcium carbonate fillers in the lumens of wood pulp fibres. The pulp is pretreated with a cationic polymer prior to being impregnated with the filler. Different conditions of pH and temperature are specified depending on whether the filler is a precipitated calcium carbonate or a ground calcium carbonate. The lumen-loaded pulps are used to make novel products with advantages in higher filler retention and sheet strength over conventionally made papers.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

METHOD FOR PRODUCING PULP AND PAPER WITH CALCIUM CARBONATE FILLER

TECHNICAL FIELD

5

This invention relates to an improved method for the production of pulps of high filler content in which filler is loaded in the lumens of the cellulose fibres and to novel pulps produced using such method.

- 10 More specifically, the present invention relates to a novel method of producing paper containing high concentrations of calcium carbonate filler and to novel paper produced by the method.

BACKGROUND ART

15

The increasing use of calcium carbonate as a filler in fine papers has been a major trend in recent years. The resultant alkaline sheets are brighter, stronger, have superior printability and are more permanent than sheets made under acidic conditions. In addition, the use of calcium carbonate is a 20 means of reducing the furnish costs by substituting fibre with less expensive filler. With these incentives, many papermakers strive to raise the filler content as high as possible. However, as filler content is increased, paper strength is reduced resulting in poor papermachine runnability. Fillers contribute nothing to paper strength themselves and 25 lower the concentration of load-bearing fibres. In addition, filler particles accumulate on exterior fibre surfaces reducing paper strength by interfering with inter-fibre bonding.

- Green, Fox and Scallan (U.S. Patent No. 4,510,020) describe one approach to improving the strength of papers containing fillers. They disclose a method of loading the filler within the fibre lumens where it does not interfere with fibre-fibre bonding. Thus, the potential is there for greater 5 filler contents in the paper and better paper machine runnability. The basic process of lumen loading involves an impregnation step in which the pulp is agitated in a concentrated suspension of filler to allow the filler particles to enter the lumens via pit apertures. If attractive forces between the filler particles and the fibre surfaces exist, the filler bonds to both exterior and 10 the lumen surfaces of the fibres. In a subsequent step the particles on the exterior surfaces of the fibres are removed by washing the pulp. For the most part, the disclosure is focused on the use of titanium dioxide fillers which proved to be very suitable for lumen loading.
- 15 Application of the lumen-loading principle to calcium carbonate fillers was mentioned as possible in U.S. Patent No 4,510,020 but no examples were given. Okayama et al. Japan Tappi, 43(5), 495, (1989) found that calcium carbonate, in the size range of commercial fillers, generally loaded to levels of less than 0.08 g/g of fibre; a much lower loading level than titanium 20 dioxide under comparable conditions. A value of 0.15 g/g was obtained on a calcium carbonate of 0.1 μm diameter - well below the size of commercially available fillers of practical and economic importance in papermaking.
- 25 Retention aids have been proposed to promote lumen loading of fillers. Middleton and Scallan, J. Pulp Paper Sci., 15(6), 229 (1989) have described the use of a cationic polyacrylamide at pH 4 to increase lumen loading using titanium dioxide. Miller and Paliwal, J. Pulp Paper Sci., 11(3), 84,

(1985) have described the use of polyethylenimine to increase the levels of lumen loading using titanium dioxide and clay fillers. A process for lumen loading calcium carbonate using polyethyleneimine is described by Chang et al, Taga Proceedings 1997 (TAGA), Session: Experimental Analysis of 5 Printing, p639-657, 1997. Using an unspecified grade of calcium carbonate, Chang et al reported loading levels of only 1-5% with a brief mention of a maximum level of 10.8% being achieved using 8% polymer addition and mixing being carried out at a pH of 13. These conditions of a very high polymer addition and a very high pH would be a severe barrier to 10 practical implementation in a mill. Another method for lumen loading calcium carbonate is reported by Hockman and Sohara, International Publication Number WO 98/35095. In this method filler and fibre are mixed together so as to effect lumen-loading. This is followed by the addition of a flocculating agent to prevent the filler diffusing outside the 15 lumens. Levels of loading of up to 10% were claimed.

There have been other approaches to producing pulps containing calcium carbonate formed by "in situ" precipitation. Allan et al. U.S. Patents Nos 5,096,539 and 5,275,699, for example, saturate fibres with calcium chloride 20 solution and then add sodium carbonate solution. However, in addition to producing calcium carbonate, the process leaves sodium chloride as a by-product which is considered detrimental in any commercial application. In an attempt to avoid such a by-product, Klungness et al. U.S. Patent No. 5,223,090 impregnate fibres with calcium hydroxide solution and then 25 apply an atmosphere of carbon dioxide to precipitate calcium carbonate. Both precipitation procedures produce calcium carbonate in various locations in a pulp. Klungness et al reported that the filler actually in the lumen was less than 0.06 g filler/g fibre.

For both "in situ" precipitation techniques, the claims in terms of benefits for the paper sheet are similar to those of lumen loading. These benefits are improved retention of filler during sheet formation and superior sheet strength over conventionally-filled sheets, i.e., where all the filler is retained on the outer surfaces of the fibres. The two precipitation techniques have common disadvantages. The first is the difficulty of obtaining an optimum size distribution of the filler for maximum optical properties. In contrast, commercial precipitated calcium carbonate is manufactured to specific particle sizes to produce optimum light-scattering characteristics. The second is that much filler is not within the lumen but external to the fibre i.e., where it causes a loss of sheet strength. In addition, "in-situ" procedures call for marked deviations from common papermaking practices.

At present, two classes of calcium carbonate fillers are commercially available. The first is a "ground" filler prepared by mechanically grinding naturally occurring deposits such as chalk or limestone. The other class is a "precipitated" filler prepared from a solution by addition of a reactant bringing about a precipitation of calcium carbonate. Within the two classes there are various grades based on particle size and shape. However, a chemical difference between the two is that the "ground" filler usually contains an adsorbed dispersant rendering its particles with a negative electrical charge while the "precipitated" filler usually has no such additive and its particles retain their natural weakly positive charge. Although the terms ground and precipitated are used in this specification it is the aspect of the electrical charge of the filler particles to which we are referring rather than the method of preparation of the filler.

DISCLOSURE OF THE INVENTION

It is an object of this invention to provide a process for production of pulp

- 5 fibres, lumen-loaded with a calcium carbonate particulate filler.

It is another object of this invention to provide pulp fibres lumen-loaded
with a calcium carbonate particulate filler.

- 10 In one aspect of the invention there is provided a process for production of
pulp fibres, lumen-loaded with a calcium carbonate particulate filler
comprising: a) contacting pulp fibres having anionically charged lumen
surfaces, with an aqueous solution of a cationic polymer with formation of
ionically charged polymer bound to the lumen surfaces, and b) contacting
15 the resultant pulp fibres with particulate calcium carbonate filler having an
ionic charge and binding the particulate calcium carbonate filler to the
lumen surfaces, such that the ionic charge on the filler is opposite to an
ionic charge on the bound polymer.
- 20 In another aspect of the invention there is provided a process for the
production of pulp fibres, lumen-loaded with a calcium carbonate
particulate filler comprising: i) agitating a suspension of pulp fibres with a
water soluble cationic polymer to form a suspension in which the pulp
fibres have the polymer bound to the lumen surfaces of the fibres, and ii)
25 adding a calcium carbonate particulate filler to the resulting suspension
from step i) and agitating so as to impregnate the lumens of the pulp fibres
with the filler.

In yet another aspect of the invention there is provided pulp fibres lumen loaded with calcium carbonate filler and having ionically charged water soluble polymer bound to the lumen surface of the fibres.

5 BRIEF DESCRIPTION OF DRAWINGS

FIG. 1 illustrates graphically the relationship between lumen-loading level achieved and temperature, employing cationic filler and hydrolysable polymer in a process of the invention;

10 FIG. 2 illustrates graphically the relationship between lumen-loading level achieved and temperature employing anionic filler and hydrolysable polymer, in a process of the invention;

15 FIG. 3 illustrates graphically the relationship between lumen-loading level and amount of polymer added;

FIG. 4 illustrates graphically the relationship between level of lumen-loading and time at different filler/fibre ratios;

20 FIG. 5 illustrates graphically the relationship between lumen-loading level and pulp consistency;

25 FIG. 6 illustrates graphically the relationship between lumen-loading level and impregnation time for different pilot plant runs; and

FIG. 7 illustrates graphically a comparison between strength properties of paper sheets formed from pulps of the invention, and paper sheets conventionally filled with cation carbonate filler.

5 DESCRIPTION OF PREFERRED EMBODIMENTS WITH REFERENCE TO THE DRAWINGS

In the first step of a particular embodiment of the method of the invention, a cationic polymeric retention aid is added to a pulp fibre suspension while 10 agitating the suspension for a period of time sufficient to cause the retention aid to enter the lumens of the fibres. In the second step, a slurry of calcium carbonate filler is added to the polymer treated fibre suspension and agitation is continued for a period of time sufficient to cause the filler to enter the lumens of the fibres, become attached to the lumen wall, and to 15 achieve sufficient loading of filler in the lumens. In an optional third step any filler attached to the external walls of the fibres may, if required, be partially or totally removed by washing the suspension.

The calcium carbonate filler is, in particular, ground calcium carbonate 20 filler having a negative charge, i.e., anionic, or precipitated calcium carbonate filler having a positive charge, i.e., cationic.

The fillers typically have a particle size of 0.4 to 1.5 μm .

25 Lumen-loadings are achieved in the order of 0.1 to 0.4 g CaCO₃/g of fibre, or 9 to 28%, by weight, of filler, based on the weight of lumen-loaded-fibres, i.e., the combined weight of the fibres, adsorbed polymer and filler in the lumen.

The polymer employed in the invention is a water soluble, cationic polymer of the type used as a retention aid to retain fillers in paper manufacture, and is employed in an aqueous solution.

- 5 The polymer is preferably a polyacrylamide containing quaternary ammonium groups attached by ester linkages to the polymer backbone and is more preferably of high molecular weight (10^5 to 10^7) and low charge density. The ester linkages are hydrolysable and thus this polymer is especially advantageous when the calcium carbonate filler has a cationic 10 charge, for example, precipitated calcium carbonate filler.

The polymer is also useful with calcium carbonate filler having an anionic charge, for example, ground calcium carbonate.

- 15 Other cationic polymers, for example, polyethylenimines, polyamines, polyamides and polydiallyldimethyl ammonium chloride, as well as cationic starch may also be used when the calcium carbonate filler has an anionic charge, for example, ground calcium carbonate filler. These 20 cationic polymers do not hydrolyse to anionic polymers and thus are less useful with cationic calcium carbonate fillers such as precipitated calcium carbonate.

In a particular embodiment of this invention, to obtain maximum lumen loading of precipitated calcium carbonate filler, the polymer is, with 25 advantage, a hydrolysable cationic polyacrylamide and the impregnation step of the process is conducted at a temperature greater than 40° C.

In yet another embodiment of the invention adapted to using ground calcium carbonates, the same process is employed but the temperature during impregnation is kept below 40°C. Alternatively a "non-hydrolysable" cationic polymer is used.

5

In a product embodiment of this invention, a pulp, the lumens of which have been loaded with calcium carbonate according to the invention, is used as part of a papermaking furnish to produce a paper which is stronger at a given filler content than a paper with all the filler conventionally-loaded onto external fibre surfaces.

10

The fibres most widely used in papermaking are cellulosic fibres derived from wood and after pulping the majority appear, under the microscope, as long hollow tubes, uniform in size for most of their length but tapered and closed at each end. Along the length of the fibre, the fibre wall is perforated by small apertures or pits which connect the central cavity or lumen to the fibre exterior. One criterion for the employment of a filler in a lumen loading process is that the filler particles are of such a size that they can enter the lumens via accessible openings, i.e., the pits or cut ends of fibres. Most commercially available calcium carbonate fillers have a particle size suitable for lumen loading.

15

A further requirement in obtaining an appreciable level of lumen loading is that there be a strong attractive force to hold the filler particle to the lumen wall. In the absence of such a force there will be no significant build-up of filler in the lumen and any small accumulation will be removed in subsequent washing or processing steps.

The lumen surface, like the exterior surface of the fibres, has a natural anionic charge due to the presence of carboxylic, and, on occasion, sulphonic acid groups within the fibre wall material. This means that ground calcium carbonates, which contain anionic dispersants conferring a negative charge to the particles, will not lumen load significantly due to the repulsive force between the like charges of filler and fibre surfaces. However, precipitated calcium carbonate, which has a small cationic surface charge will be retained by the fibres to some degree due to the small attractive force between the cationic filler and the anionically-charged lumen-surface. Nevertheless, even with precipitated calcium carbonate, the level of lumen-loading is still too low to be practically useful.

In the process of the invention, to achieve high levels of lumen loading with calcium carbonate in excess of 9%, by weight, pulp fibres dispersed in water as a suspension, are first treated with a cationic polymer and agitation is employed to cause the polymer to be adsorbed on the exterior and lumen surfaces of the pulp fibres. Five minutes of agitation is found to be sufficient. Due to its cationic charge, the polymer readily adsorbs onto the anionic fibre surfaces. Following the polymer addition, precipitated calcium carbonate filler, pre-dispersed as a suspension in water usually at 20% solids, is added and the fibres are impregnated with filler using vigorous agitation. During impregnation, the filler enters the lumens and attractive colloidal forces, induced by the polymer, hold the filler particles onto the lumen wall. Following completion of the impregnation step a significant fraction of the filler remains free in suspension and on the external walls of the fibres. Optionally, the fibres can be made substantially free of external filler by washing the pulp while containing it

by a screen which will permit passage of filler particles but not the fibres. Sufficient shear is introduced during the washing action to overcome the attractive colloidal forces holding the filler particles to the external surface but not to unduly dislodge particles in the lumens. The particles in the 5 lumen are protected to some extent from the shear forces by the fibre wall although some loss of this filler will occur. For this reason, it is preferable not to prolong washing beyond the time necessary to remove the external filler on the fibres.

- 10 In a further embodiment of the invention applied to achieving high levels of lumen-loading of precipitated calcium carbonate fillers, the polymer of choice is a cationic polyacrylamide polymer such as Percol 292 (Trademark of Allied Colloids Inc.). Preferably, the polymer pretreatment is carried out with the fibre suspension below pH 7 and the impregnation step 15 is carried out at an elevated temperature, preferably greater than 40°C and at a pH greater than 8. The alkaline pH value is achieved naturally by the addition of the calcium carbonate. A graphical illustration showing the preferred embodiment of elevated temperature to achieve high loading levels when using precipitated calcium carbonate is shown in Fig. 1. These 20 surprising results are believed to be due to the effect of pH and temperature on the cationic polyacrylamide. The cationicity of the polymer arises from quaternary ammonium groups attached by ester linkages to the polymer backbone. Under conditions of alkalinity and accelerated by heat, hydrolysis of the ester linkages occurs, the polymer loses its cationic charge 25 and gains an anionic charge arising from the acid groups formed on the polymer as residuals of the ester linkages. Thus the initial cationicity of the polymer achieves adsorption of the polymer onto the negatively charged fibre lumen surfaces. When the precipitated calcium carbonate is added to

start impregnation, the pH of the suspension naturally becomes alkaline and if the suspension is heated to 40°C or more, hydrolysis occurs. However, in spite of the charge reversal, the polymer still remains attached to the fibre lumen wall and the anionic charge produced on the polymer favours attachment of the cationic precipitated calcium carbonate.

It has been found that, when ground calcium carbonate is lumen loaded using the same polymer and the same physical conditions as for the preceding experiments, higher impregnation temperatures are detrimental.

10 This is probably because hydrolysis of the polymer leaves both fibre surface and filler negatively charged. A further embodiment of the invention for anionic fillers coupled with hydrolysable polymers, is that a high loading level is favoured by operating the impregnation step at temperatures less than 40°C as illustrated in Fig. 2 so as to avoid hydrolysis

15 and maintain the cationic charge on the polymer. Alternatively, in yet a further embodiment using ground calcium carbonate filler, a non-hydrolysable cationic polymer can be employed and then the impregnation temperature is immaterial. As indicated above the polymer can be chosen from a large group of cationic polymers currently used in papermaking

20 furnishes, including cationic starch, polyethylenimine, polyDADMAC (polydiallyldimethyl ammonium chloride), polyamine and polyamide.

The foregoing features of the invention have been disclosed to show when and how certain calcium carbonate fillers can be appreciably loaded or not

25 and these are the primary features of the invention. Additionally, several other variables affecting the actual level of loading have been discovered. The effect of the level of addition of polymer is illustrated in Fig. 3. An

addition of 0.05% of Percol 292 (Trade-mark) is indicated as producing an observable effect but a 0.5% addition is preferred.

The effect of time of impregnation is illustrated in Fig. 4. While
5 appreciable loading is achieved in 20 mins, 60-120 mins is preferred in order to effect maximum loading. Fig. 4 also illustrates that the higher the filler to fibre ratio, the higher the level of lumen loading. Very high ratios are not too practical to achieve high loading and in general a weight ratio of filler to fiber of 0.5:1 to 3:1 is employed in the process.

10 The results in Fig. 5 illustrate how, at a low filler to fibre ratio, loading may be greatly increased by employing higher pulp consistencies. It is believed that the reason for this phenomenon is that the driving force for loading is the concentration of filler in suspension rather than the filler to fibre ratio
15 per se. From these findings it will now be clear to a person skilled in chemical kinetics how one may obtain optimum performance with the combinations of novel variables at his disposal.

20 Experimentation has been on a laboratory scale using thirty grams of pulp per run, but the preferred procedure has also been demonstrated for precipitated calcium carbonate carried out in a pilot plant handling 27 kg of pulp per run. Notably the procedure is translatable to the larger scale without change in kinetics, thus the time scale of impregnation is the same. Four separate runs with the same set of conditions showed the procedure to
25 be highly reproducible. An inclined screen device was shown to be a practical means for washing on a large scale.

As to the use of lumen-loaded pulp made according to the novel process of the invention, the washed product can be used as part of a furnish containing other pulps, additives and fillers. The advantage of adding lumen-loaded pulp as a component to the furnish is that the filler contained 5 within the fibre will have less of a weakening effect on the sheet than externally held filler as shown in Fig. 7. This aspect may be utilized to increase the filler content of the paper sheet or alternatively to obtain a stronger and better running sheet at the same filler content.

- 10 In an alternative application of the lumen-loaded pulp, the loaded pulp after the impregnation step is not washed free of the unloaded filler but mixed directly with other stock in the papermaking furnish. One example would be a fine paper mill using a softwood/hardwood furnish and producing a sheet containing calcium carbonate. In such an application it would be 15 advantageous to use all the filler intended for the final sheet, in a lumen loading treatment of the softwood fibres thus confining treatment to the most responsive pulp and obtaining the high filler to fibre ratio necessary for high loading. Following loading and without washing, the hardwood pulp could then be added to the furnish. Although the final sheet will 20 contain a large fraction of filler conventionally-loaded, a significant fraction of the loading will be in the lumen bringing some benefits in terms of retention, sheet strength and hence runnability. These factors will permit a higher level of filler in the sheet and hence a reduction in furnish cost.
- 25 Pulp fibres lumen-loaded with calcium carbonate made according to this invention can be used in a wide variety of applications including fine papers, light-weight newsprint, newsprint specialities etc. Without further elaboration, it is believed that one skilled in the art can, using the preceding

description, utilize the present invention to its fullest extent. The following are further illustrations of the novel findings and should not limit the scope of this invention in any way.

5 EXAMPLES

EXAMPLE 1

30 g dry weight of a bleached never-dried softwood kraft pulp was diluted 10 to 1000 g with deionised water and dispersed in a mixing device (British disintegrator) for 5 mins at 3000 rpm. To this suspension, a cationic polyacrylamide (Percol 292 from Allied Colloids Inc.) was added to give 0.5% by weight on pulp. The polymer was added as a 1 g/L solution previously prepared from dry polymer by gentle stirring in deionised water 15 for 24 hours. Adsorption onto the pulp was allowed to occur during 10 min stirring at 1000 rpm. Then, 90 g (dry weight) of a precipitated calcium carbonate filler (Albafil M Trade-mark of Specialty Minerals Inc.), predispersed in water at 20% concentration, was added to the pulp. Finally, sufficient water was added to raise the total weight of water in the British 20 disintegrator to 1500 g. The mixture was then stirred for 1 hour at 1000 rpm at a temperature of 75°C to effect loading.

Following the impregnation, the fibre/filler mixture was washed in tap water (8L/min) in a single unit of a Bauer-McNett classifier (equipped with 25 a 100 mesh screen) until the fibre was free of external filler (10 min). The filler content within the lumens was calculated from the ash content of the pulp determined at 900°C and was found to be 0.28 g filler/g fibre.

EXAMPLE 2

The same procedure to that described in Example 1 was repeated but with
5 the impregnation step carried out at a series of temperatures between 25
and 75°C. The ratio of filler to fibre was 2:1 and the impregnation time
was 20 minutes, otherwise conditions were as for Example 1. Fig. 1 shows
the results of these experiments and illustrates the beneficial effect of
temperature in the case of precipitated calcium carbonate filler.

10

EXAMPLE 3

The procedure given in Example 1 was carried out on a series of different
15 pulp fibres. As shown in Table 1, all the pulps respond to the lumen-
loading treatment but there is a variation in loading level due to the nature
of the fibres.

Table 1 - Calcium carbonate loading (g filler/g fibre) as a function of pulp type.

Bleached kraft, softwood, never dried, unbeaten	0.28
Bleached kraft, softwood, never dried, beaten	0.27
Bleached kraft, softwood, dry-lap, rewetted	0.17
Unbleached kraft, softwood, never dried	0.27
Thermomechanical, softwood	0.23

5

EXAMPLE 4

The procedure given in Example 1 was carried out on a scalenohedral type
 10 of precipitated calcium carbonate of particle size 1.3 μm . The loading level
 was 0.14 g filler/g fibre compared to that of 0.28 g/g for the smaller filler
 (size 0.8 μm) cited in Example 1.

EXAMPLE 5

15

The procedure given in Example 2 was repeated using a ground calcium carbonate as filler (Omyafil from Omya Inc.). The level of lumen-loading
 was 0.17 g filler / g fibre at 25°C falling to 0.02 g filler/g fibre at 75°C.
 This result is given in Fig. 2 showing a preferred embodiment of an
 20 impregnation temperature less than 50°C when a hydrolysable polymer and

a ground calcium carbonate are used. An impregnation temperature of less than 40°C is yet more preferable in improving the level of lumen loading.

EXAMPLE 6

5

The procedure given in Example 1 was carried out with a 2:1 ratio of filler to fibre, an impregnation time of 20 minutes, and a series of different polymer addition levels. Fig. 3 illustrates the effect of polymer addition on the level of lumen-loading showing a preferred embodiment of a cationic 10 polyacrylamide polymer addition of at least 0.1% on pulp.

EXAMPLE 7

15

The procedure given in Example 1 was carried out a number of times and variations were made in impregnation time and the filler to fibre ratio. Increases in both these parameters result in higher levels of lumen loading as illustrated in Fig. 4. Thus, preferred embodiments to achieve high levels of lumen loading, are the use of high filler to fibre ratios and, at any given 20 filler to fibre ratio, extending the time of impregnation until a maximum in loading is obtained.

EXAMPLE 8

A series of lumen-loading procedures were carried out at various pulp 25 consistencies following the procedure given in Example 1 except that the filler to fibre ratio was held at 1:1 and the mixing speed during impregnation was 2000 rpm. The results of these experiments are given in Fig. 5 and illustrate that, to achieve high levels of lumen loading at any

given filler to fibre ratio and impregnation time, a preferred embodiment is the use of as high a pulp consistency as possible in the impregnation stage.

EXAMPLE 9

5

A pilot plant for producing lumen-loading pulp was assembled and four runs were made. A never-dried bleached kraft pulp made from softwoods was used for these runs. The polymer was a cationic polyacrylamide (Percol 292, Allied Colloids Inc.) and, prior to each run, 200 g of dry polymer was gently stirred in 200 L of deionized water for 16 hrs at 25°C. 10 The filler was a dry precipitated calcium carbonate (Albafil M, Specialty Minerals Inc.) and, prior to each run, 54 kg was dispersed in 162 kg tap water using a Cowles mixer.

15 To start a run, 27 kg of the pulp with associated water (total weight 163 kg) was added to 700 L tap water at 96°C in a baffled tank of capacity 3000 L. The pulp was stirred at 300 rpm with a 4-blade rotor for 5 minutes to achieve good dispersion. The polymer solution was then added to the pulp (0.75% polymer on pulp) and mixing carried out for 10 min at 300 rpm and 20 a temperature of 80°C.

The filler suspension was then added to the pulp (giving a 2:1 filler:fibre ratio) and impregnation was carried out at 300 rpm for 3 hours at 60°C. During the impregnation, samples of pulp were taken from the suspension 25 and the degree of lumen-loading determined by washing and ashing, as in Example 1. In Fig. 6 are shown the results from four separate but similar runs and it is seen that the loading initially increases rapidly with time but

then reaches a plateau value within 1 hour. After 3 hours the agitation was stopped.

Following impregnation, the pulp was washed. This was accomplished by
5 diluting the stock with tap water to 0.5% consistency and then pumping it over a Sidehill-type screen washer to separate fibre and filler. After this the pulp, now at 5% consistency, was diluted with fresh tap water. The washing process was carried out for a total of 4 cycles. After the final washing the loaded pulp was pressed to a consistency of 20%. It was
10 determined that this material had 0.25 g filler/g fibre within the lumens and 0.05 g filler/g fibre on external surfaces.

This example serves to show that the laboratory procedure is translatable to a pilot scale and the inference is that the process could be further scaled up
15 to an industrial level.

EXAMPLE 10

Example 1 was repeated several times but varying the filler to fibre ratio
20 and the impregnation time to produce a series of pulps lumen-loaded to different degrees. These pulps were made into handsheets and the tensile strength properties of the sheets measured. In Fig. 7 we show a comparison of the strength properties of these handsheets with handsheets made with the same filler retained conventionally (i.e., on external fibre surfaces).
25 These results illustrate that, an advantage of paper lumen-loaded with calcium carbonate filler over paper conventionally loaded with the same filler, is superior tensile strength at any given filler content.

EXAMPLE 11

The procedure given in Example 2 was repeated but using a precipitated calcium carbonate which had been treated with 0.5% tetrasodium pyrophosphate, an anionic dispersant. When impregnation is carried out at 5 25°C, the lumen-loading level was 0.17 g filler/g fibre but when impregnation was carried out at 75°C, the lumen-loading level dropped to 0.01 g/g. This illustrates the preferred embodiment of temperatures below 40°C when the calcium carbonate filler is anionic, irrespective of whether it 10 is ground or precipitated.

CLAIMS

1. A process for production of pulp fibres, lumen-loaded with a calcium carbonate particulate filler comprising:
 - 5 a) contacting pulp fibres having anionically charged lumen surfaces, with an aqueous solution of a cationic polymer with formation of ionically charged polymer bound to the lumen surfaces, and
 - b) contacting the resultant pulp fibres with particulate calcium carbonate filler having an ionic charge, and binding the particulate calcium carbonate filler to the lumen surfaces, under conditions such that the ionic charge on the filler is opposite to an ionic charge on the bound polymer.
- 10 2. A process according to claim 1, wherein in step a) pulp fibres are formed having cationically charged polymer bound to the lumen surfaces, and the particulate calcium carbonate filler in step b) is a ground calcium carbonate filler having an anionic charge.
- 15 3. A process according to claim 1, wherein said cationic polymer in step a) has hydrolysable ester groups and the particulate calcium carbonate filler in step b) is precipitated calcium carbonate filler having a cationic charge.
- 20 4. A process according to claim 1, 2 or 3, wherein said cationic polymer in step a) is a polymeric retention aid for filler loading of pulp in paper manufacture.

5. A process according to claim 1, comprising:

- i) agitating a suspension of the pulp fibres with a water soluble cationic polymer to form a suspension in which the pulp fibres have the polymer bound to the lumen surfaces of the fibres, and
- 5 ii) adding a calcium carbonate particulate filler to the resulting suspension in step i) and agitating to impregnate the lumens of the pulp fibres with the filler.

10 6. A process according to claim 5, further including:

- iii) washing the pulp fibres from step ii) to remove filler from external surfaces of the fibres.

7. A process according to claim 5 or 6, wherein said cationic polymer
15 in step i) comprises a copolymer of acrylamide and acrylic acid monomers, said copolymer bearing quaternary ammonium groups attached by ester linkages to acid groups of the copolymer, said ester linkages being hydrolysable, and said quaternary ammonium groups rendering said polymer cationic.

20

8. A process according to claim 5, 6 or 7, wherein said agitating in step i) is under acid pH whereby said cationic polymer is adsorbed on the lumen surfaces of the pulp fibres; the filler in step ii) is cationic and step ii) is carried out under alkaline conditions and at a temperature effective to 25 hydrolyse said ester linkages to render the adsorbed polymer anionic.

9. A process according to claim 8, wherein said filler is precipitated calcium carbonate filler.

10. A process according to claim 5, wherein said filler has been pretreated with cationic dispersant or cationic polymer.
- 5 11. A process according to claim 5, 6, 7, 8 or 9, wherein said polymer in step i) is present in an amount of 0.01% to 1.0%, by weight, based on the oven dry weight of the pulp fibres.
- 10 12. A process according to claim 5, 6 or 7, wherein said filler in step ii) is anionic, and steps i) and ii) are carried out under conditions of temperature and pH such that said ester linkages are maintained non-hydrolysed.
- 15 13. A process according to claim 12, wherein said filler is ground calcium carbonate filler.
14. A process according to claim 12, wherein said filler is precipitated calcium carbonate filler rendered anionic by pre-treatment with an anionic dispersant or polymer.
- 20 15. A process according to claim 5, wherein said cationic polymer is selected from polyamine, polyethylenimine, poly DADMAC, polyamide and cationic starch and said filler is anionic.
- 25 16. A process according to any one of claims 1 to 15, wherein the pulp fibres from step b) together with filler on the external surface of the fibres is added directly to a paper furnish.

17. A process according to any one of claims 1 to 16, wherein the lumen-loaded fibres from step b) have a lumen content of calcium carbonate filler of 9 to 28%, by weight, based on the weight of the lumen-loaded fibres.

5

18. A process according to any one of claims 1 to 11, wherein said polymer in step i) is of a polymeric retention aid having a weight average molecular weight of 1×10^5 to 1×10^7 for filler loading of pulp in paper manufacture.

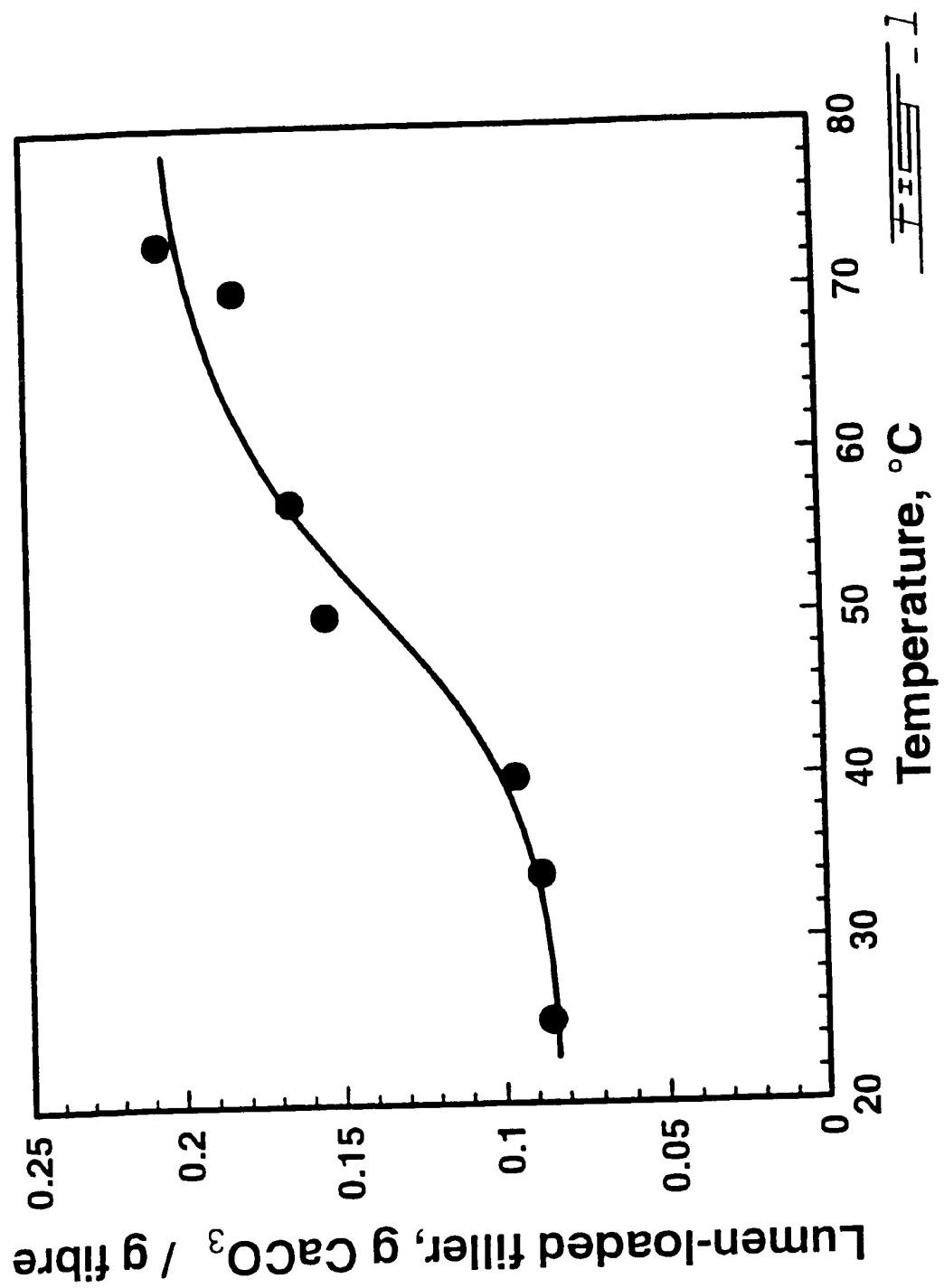
10

19. Pulp fibres lumen loaded with calcium carbonate filler and having ionically charged water soluble polymer bound to the lumen surface of the fibres.

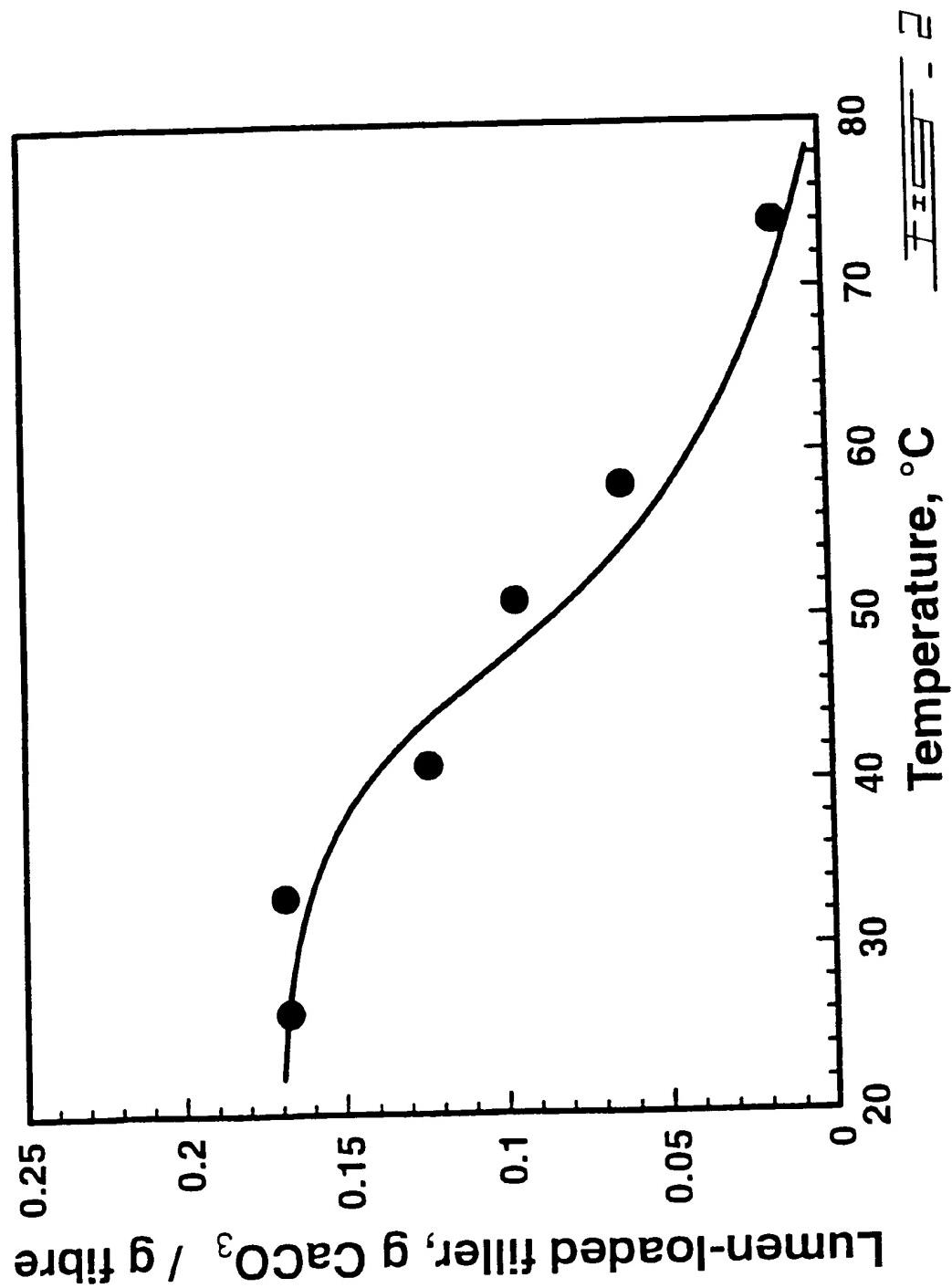
15

20. Pulp fibres according to claim 19, wherein the lumen content of calcium carbonate filler is 9 to 28%, by weight, based on the weight of the lumen-loaded fibres.

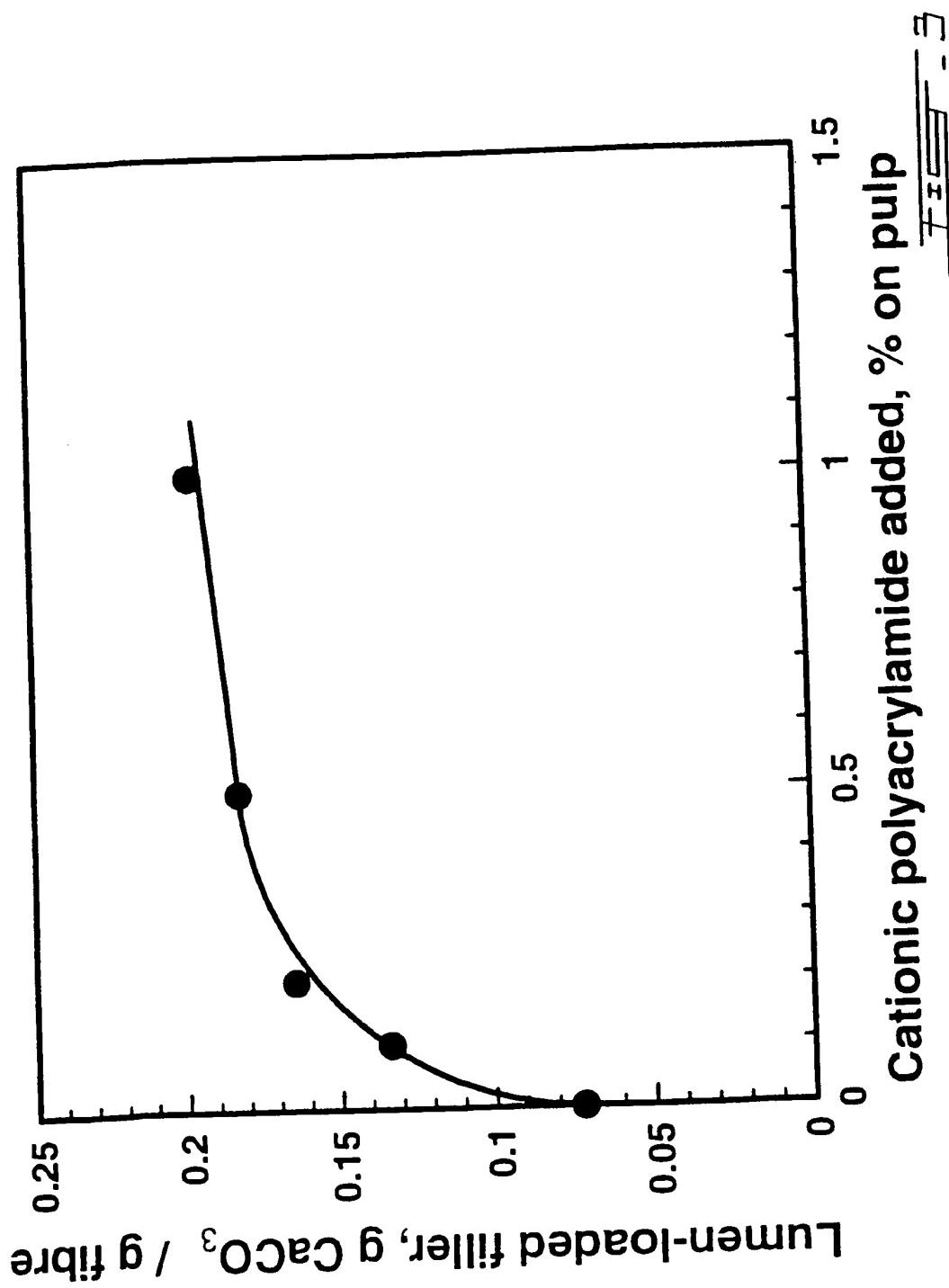
1/7



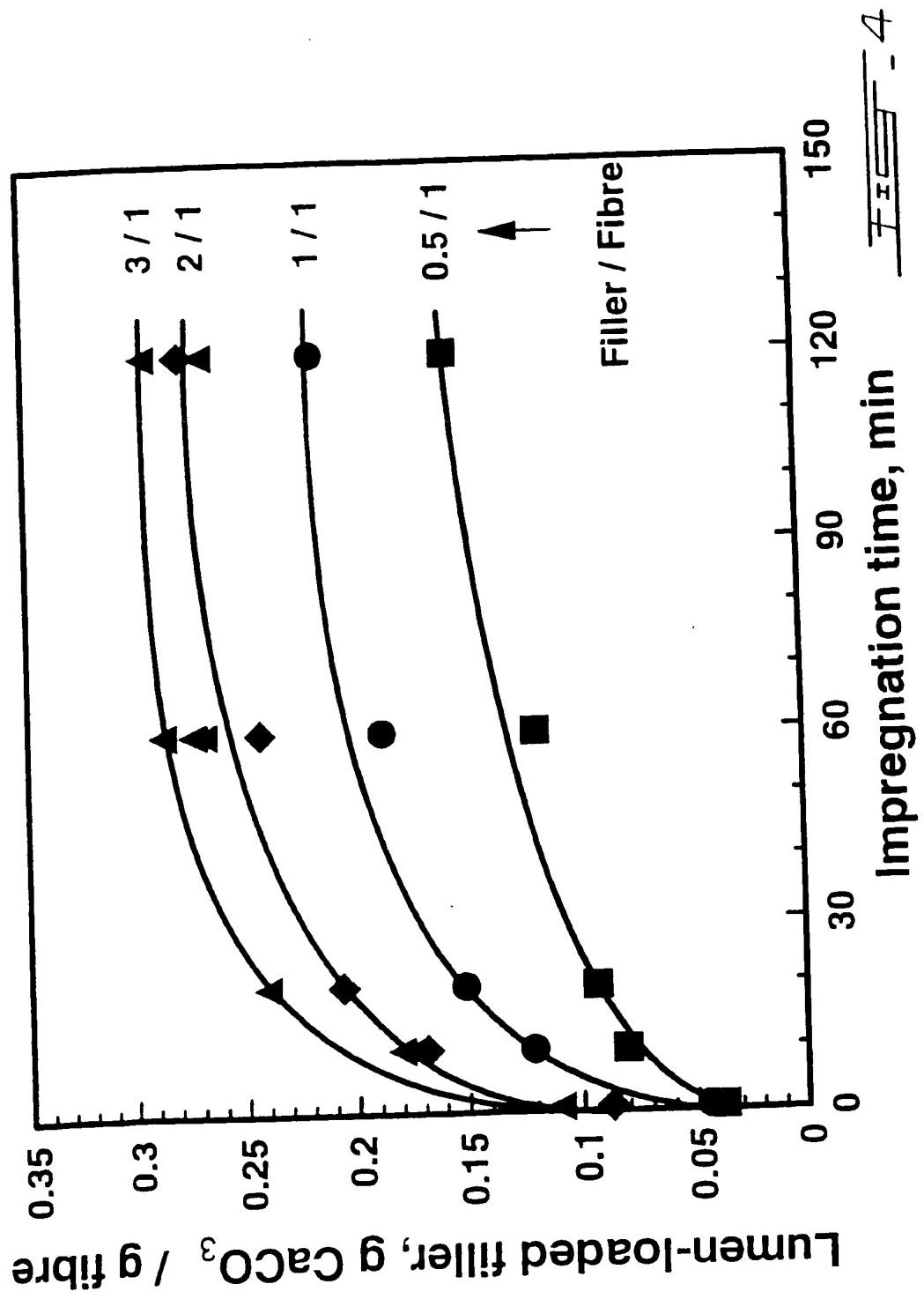
2/7



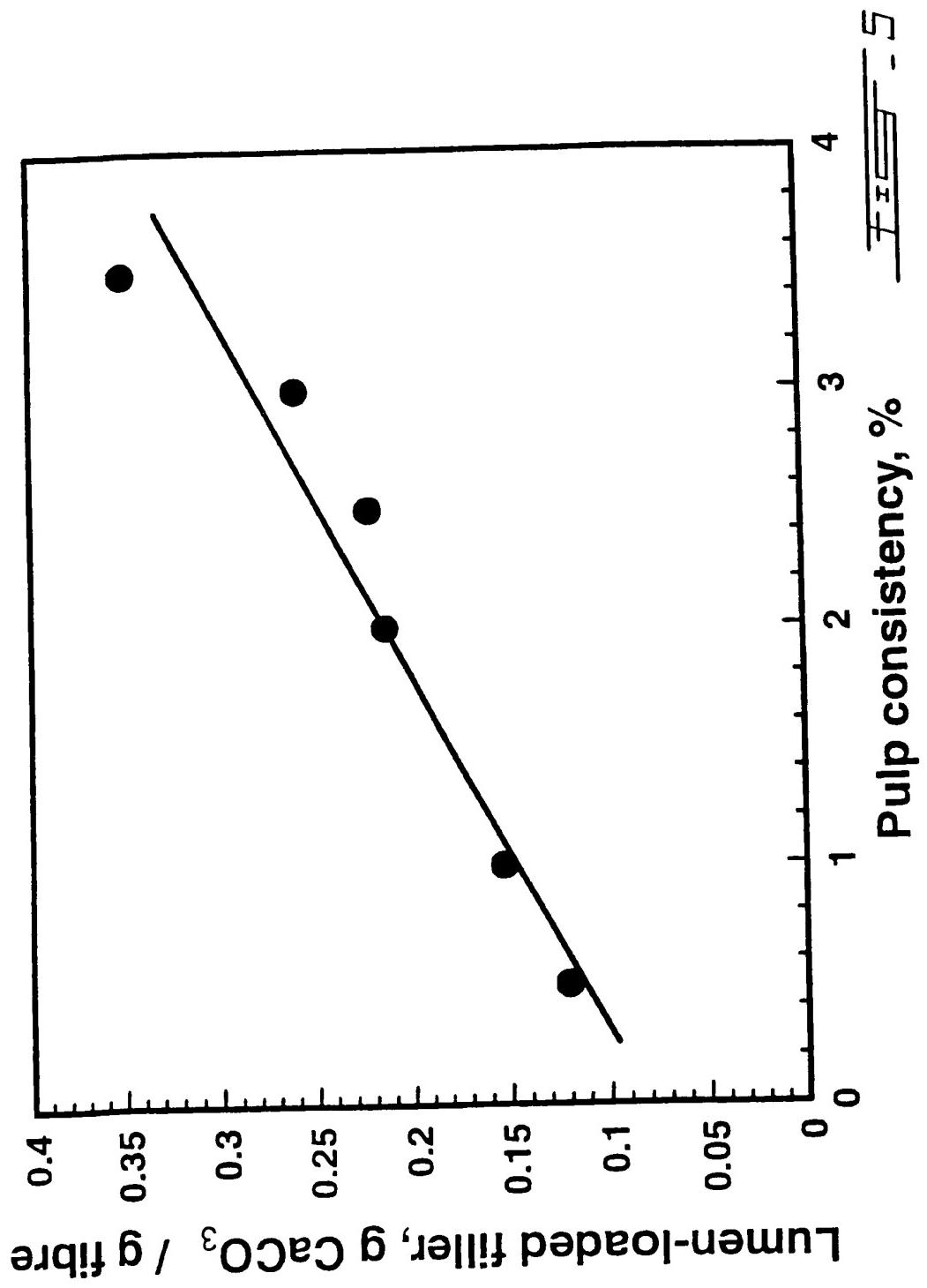
3/7



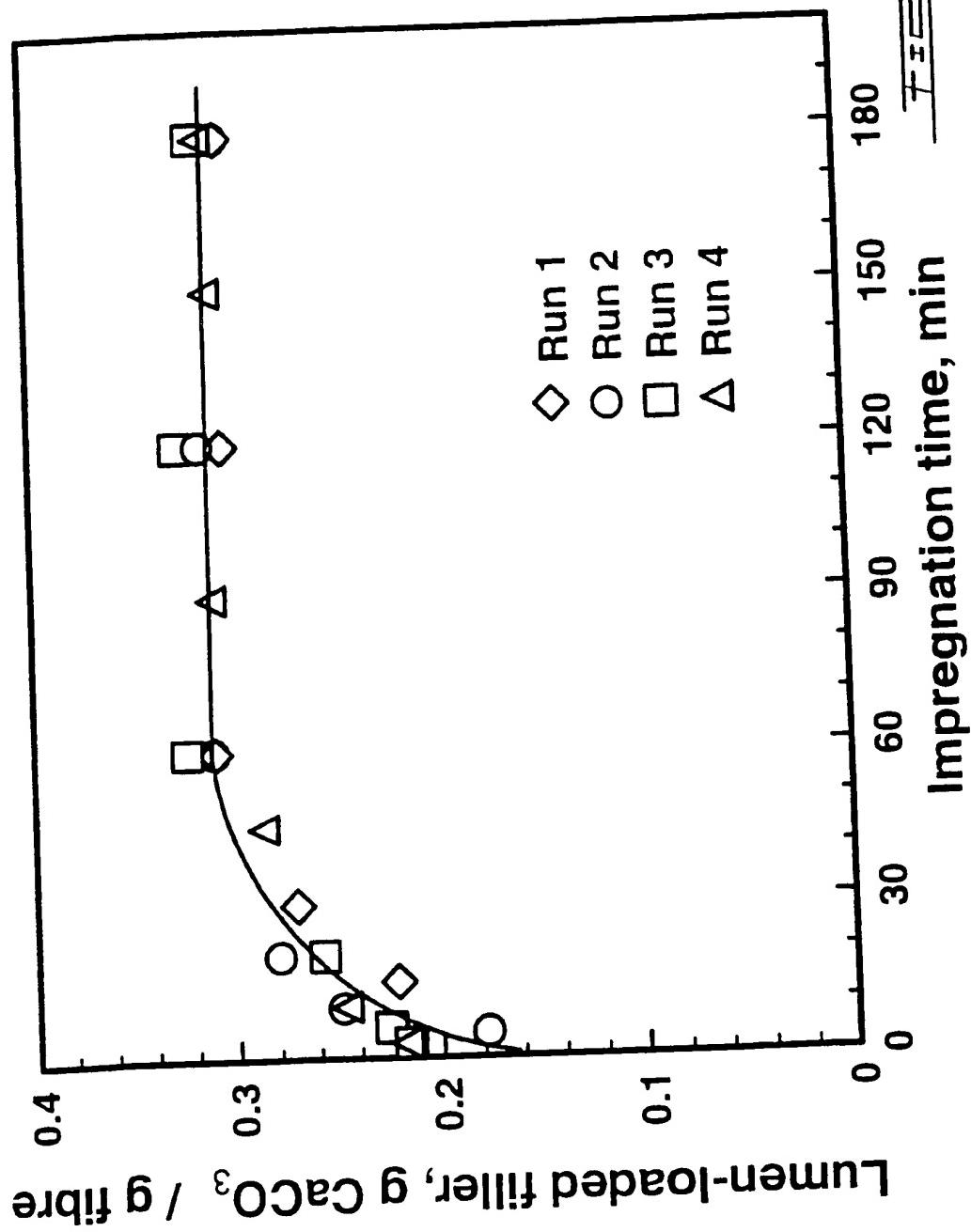
4/7



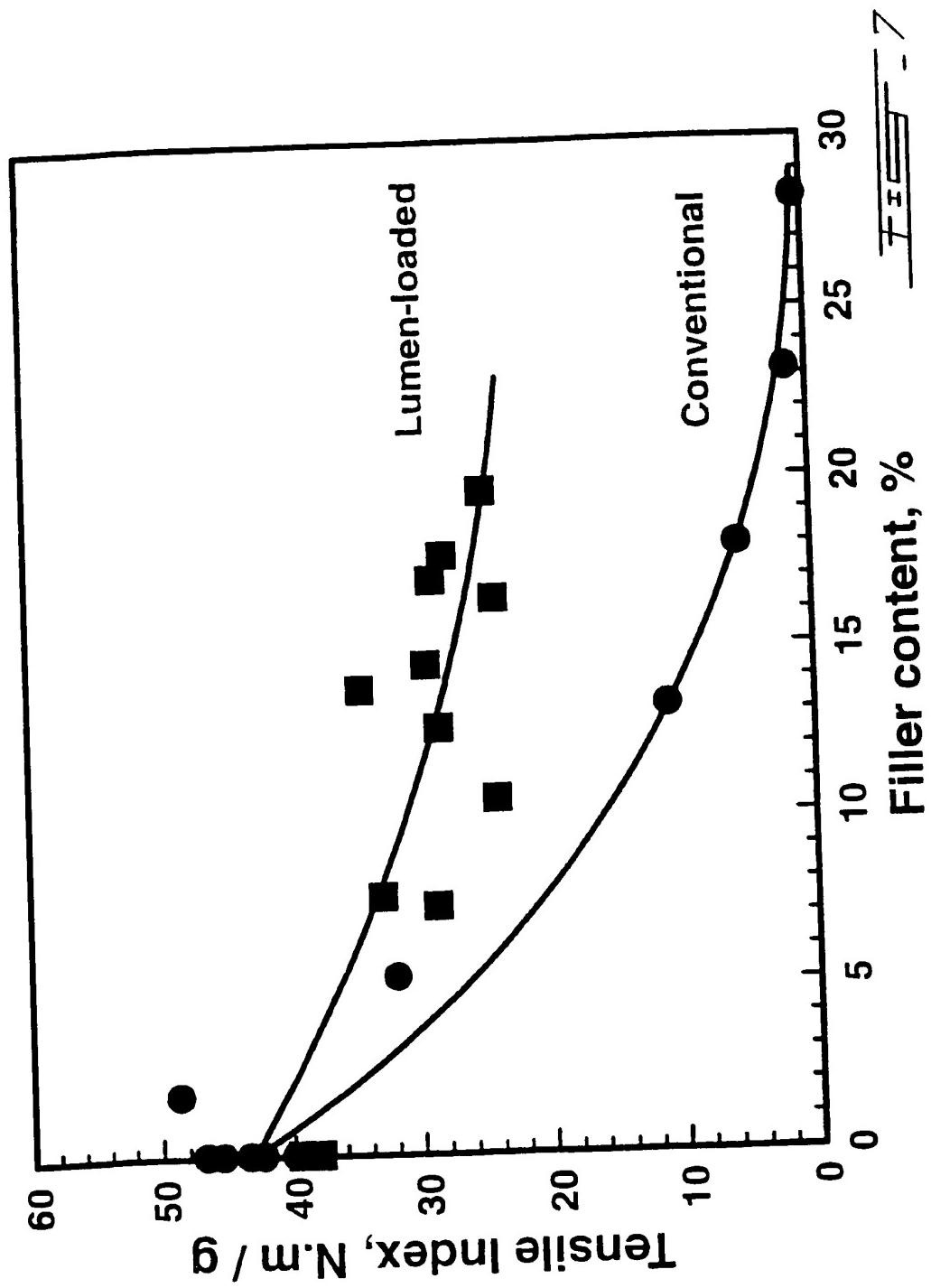
5/7



6/7



7/7



INTERNATIONAL SEARCH REPORT

International Application No
PCT/CA 99/00213

A. CLASSIFICATION OF SUBJECT MATTER

D 21 H 17/67, D 21 H 17/64

According to International Patent Classification (IPC) or to national classification and IPC6

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

D 21 H 17/00

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5679220 A (MATTHEW et al.) 21 October 1997 (21.10.97), claims 1-10. --	1-20
A	EP 0491346 A1 (HERCULES INCORPORATED) 24 June 1992 (24.06.92), claim 1. --	1-20
A	WO 97/37081 A1 (THE PROCTER & GAMBLE COMPANY) 09 October 1997 (09.10.97), claims 1-10. --	1-20
A	WO 98/13549 A1 (THE PROCTER AND GAMBLE COMPANY) 02 April 1998	1-20

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- 'A' document defining the general state of the art which is not considered to be of particular relevance
- 'E' earlier document but published on or after the international filing date
- 'L' document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- 'O' document referring to an oral disclosure, use, exhibition or other means
- 'P' document published prior to the international filing date but later than the priority date claimed

'T' later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

'X' document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

'Y' document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

'&' document member of the same patent family

Date of the actual completion of the international search

02 June 1999

Date of mailing of the international search report

07.07.1999

Name and mailing address of the ISA
European Patent Office, P.O. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+ 31-70) 340-2040, Tx. 31 651 epo nl.
Fax (+ 31-70) 340-3016

Authorized officer

BRUS e.h.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/CA 99/00213

-2-

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
	(02.04.98), claims 1-10. -----	

ANHANG

zum internationalen Recherchenbericht über die internationale Patentanmeldung Nr.

ANNEX

to the International Search Report to the International Patent Application No.

ANNEXE

au rapport de recherche international relatif à la demande de brevet international n°

PCT/CA 99/00213 SAE 226776

In diesem Anhang sind die Mitglieder der Patentfamilien der im obengenannten internationalen Recherchenbericht angeführten Patentdokumente angegeben. Diese Angaben dienen nur zur Orientierung und erfolgen ohne Gewähr.

This Annex lists the patent family members relating to the patent documents cited in the above-mentioned International search report. The Office is in no way liable for these particulars which are given merely for the purpose of information.

La présente annexe indique les membres de la famille de brevets relatifs aux documents de brevets cités dans le rapport de recherche International visé ci-dessus. Les renseignements fournis sont donnés à titre indicatif et n'engagent pas la responsabilité de l'Office.

Im Recherchenbericht angeführtes Patentdokument Patent document cited In search report Document de brevet cité dans le rapport de recherche	Datum der Veröffentlichung Publication date Date de publication	Mitglied(er) der Patentfamilie Patent family member(s) Membre(s) de la famille de brevets	Datum der Veröffentlichung Publication date Date de publication
US A 5679220	21-10-1997	keine - none - rien	
EP A1 491346	24-06-1992	AT E 124740 AU A1 89742/91 AU B2 642061 BR A 9105473 CA AA 2057548 DE CO 69111034 DE T2 69111034 EP B1 491346 ES T3 2076452 FI A0 915894 FI A 915894 JP A2 4339867 MX A1 9102608 NO AD 914953 NO A 914953 US A 5169441 ZA A 9109904	15-07-1995 18-06-1992 07-10-1993 01-09-1993 18-06-1992 10-08-1995 21-12-1995 05-07-1995 01-11-1995 13-12-1991 18-06-1992 26-11-1993 10-08-1992 16-12-1991 18-06-1992 08-12-1992 30-09-1992
WO A1 9737081	09-10-1997	AU A1 24550/97 EP A1 891444 US A 5700352	22-10-1997 20-01-1999 23-12-1997
WO A1 9813549	02-04-1998	AU A1 48015/97 US A 5759346	17-04-1998 02-06-1998